51	PENNFO	RM	ATION DISC	LOSURE		ATTY. DOCI 114232.104	KET NO.	SERIAL NO. 09/518,081	
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	EXAMINER'S INITIALS		PATENT NO.	DATE		NAME	CLASS	SUBCLASS	FILING DATE
	/W.M	/AA	4,021,472	5/3/77	Fuji		/	BOBCLINGS	DATE
	1	AB	4,051,842	10/4/977		el et al.	/	· ·	
		AC	4,140,122	2/20/79		l et al.		1	
		AD	4,224,342	9/23/80	Fuji	et al			·
		AE	4,283,418	8/11/81	_	et al			
		AF	4,310,533	1/12/82	Ueg	ai eț∕al			
		AG	4,383,529	5/17/83	Web	ster			
		AH	4,629,567	12/16/86	Boli	en et al.			
		ΑI	4,668,504	5/26/87	Kah	an et al.			
		AJ	4,711,848	12/8/87	Ínsle	ey et al.			
	мання	AK	4,713,224	12/15/87	Tam	hankar et al.			
	***************************************	AL	4,732,973	3/22/88/	Bar	r et al			
		AM	4,760,130	7/26/88	Tho	mpson et al.			
		AN	4,788,603	11/29/88	Fuji	mura et al.			
	$\rightarrow$ $\perp$	AO	4,829,052	5/9/89	Glo	ver et al.		1	
		AP	4,839,283	6/13/89	Kaw	asaki et al			
_		AQ	4,843,094	6/27/89	Imal	ki et al			
y	ľ	AR	4,889,723	12/26/89	Kim	et al			
P		AS	4,931,279	6/5/90	Baw	a et al.			
		ΑT	4,963,654	10/16/90	Katı	inuma			
		AU	5,004,612	4/2/91	Kim	et al			
		ΑV	5,077,428	12/31/91	Ima	ki et al			
		AW/	5,110,602	5/5/92	Kim	et al			
		ĄX	5,157,019	10/23/92	Glo	ver et al.			
		AY	5,175,253	12/29/92		on et al.			
		ΑZ	5,214,191	5/25/93	Kirsc	henheuter et al			
		AAA	5,240,956	8/31/93	Kirsc	henheuter et al			
		AAB	5,247,084	9/21/93	Ima	ki et al			
	/ /W.M	AAC	5,281,617	1/25/94	Kirsc	henheuter et al			
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ATTY. DOCKET NO. 114232.104 SERIAL NO. 09/518,081

APPLICANT

Leland SHAPIRO

FILING DATE GROUP March 3, 2000 1646

EXAMINER'S		PATENT NO.	DATE	NAME	CLASS	SUBCLASS	FILING
INTIALS	AAD	5,314,910	5/24/94	Kirschenheuter et al	CLASS	SOBCLASS	DAIL
	AAE	5,376,655	12/27/94	Imaki et al			
	AAF	5,399,346	3/21/95	Anderson et al.			,
-	AAG	5,416,191	5/16/95	Cheronis et al			
	AAH	5,420,110	5/30/95	Miller et al.			
	AAI	5,432,178	7/11/95	Nakai et al			
	AAJ	5,470,970	11/28/95	Saeger et al.			
	AAK	5,476,995	12/19/95	Clark et al			
	AAL	5,478,727	12/26/95	Roizman et al			
	AA M	5,486,470	1/23/96	Darke et al			
	AAN	5,514,653	5/7/96	Perlmutter			
	AAO	5,514,713	5/7/96	Nakai et al			
	AAP	5,529,920	6/25/96	Cole et al.			
	AAQ	5,532,215	7/2/96	Lezdey et al.			
	AAR	5,565,334	10/15/96	Kufe et al			
	AAS	5,593,858	1/14/97	Fleer et al			
"	AAT	5,604,201	2/18/97	Thomas et al.			
	AAU	5,610,140	3/11/97	Goodfellow et al			
	AAV	5,610,285	3/11/97	Lebing et al.			
	AA W	5,614,555	3/25/97	Nakai et al			
	AAX	5,616,693	4/1/97	Hwang et al.			
	AAY	5,618,792	4/8/97	Gyorkos et al.			
	AAZ	5,622,984	4/22/97	Nakai et al			
	ABA	5,635,593	6/3/97	Cheronis et al			
	ABB	5,641,670	6/24/97	Treco et al			
	ABC	5,663,416	9/2/97	Kirschenheuter et al			

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ATTY. DOCKET NO. 114232.104 SERIAL NO. 09/518,081

APPLICANT Leland SHAPIRO

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INITIALS	ABD	PATENT NO. 5,665,589	9/9/97	NAME Harris et al	CLASS	SUBCLASS	DATE
	ABE						
	$\vdash$	5,700,779	12/23/97	Goodfellow et al			
	ABF	5,710,026	1/20/98	Sprecher			
	ABG	5,712,117	1/27/98	Sprecher			
	ABH	5,747,645	6/6/98	Sprecher			
	AJt	5,750,506	5/12/98	Goodfellow et al			
	ABJ	5,759,548	6/2/98	Bathurst et al.			
	ABK	5,780,009	7/14/98	Karatzas et al			
	ABL	5,798,442	8/25/98	Gallant et al.			
	AB M	5,801,148	9/1/??	Gyorkos et al			
	ABN	5,807,829	9/15/98	Gyorkos et al			
	ABO	5,811,241	9/22/98	Goodfellow et al			
	ABP	5,817,484	10/6/98	Yu et al			
	ABQ	5,834,431	11/10/98	Stewart et al			
	ABR	5,843,900	12/1/98	Cheronis et al			
	ABS	5,849,863	12/15/98	Stewart et al			
	ABT	5,861,299	1/19/99	Archibald et al			
	ABU	5,861,380	1/19/99	Gyorkos et al.			
	ABV	5,863,899	1/26/99	Cheronis et al			
	AB W	5,869,455	2/9/99	Gyorkos et al.			
	ABX	5,874,424	2/23/99	Batchelor et al.			
	ABY	5,891,852	4/6/99	Gyorkos et al			
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,	ВВ	WO 98/20034	5/14/98	PCT					
	BC	WO 98/23565	6/4/98	PC1				<u> </u>	_
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	BF	WO 97/21690	6/19/97	PCT					
	BG	WO 97/10231	3/20/97	PCT					
	вн	WO 97/03679	2/6/97	PCT					
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•	вк	WO 97/48706	12/24/97	PCT	•			1	
	BL	WO 97/33996	9/18/97	PCT	•				
	ВМ	WO 98/46597	10/22/98	PCT					
·	BN	WO 97/24339	7/10/97	PCT				Abstr	
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	BQ	WO95/28422	10/26/95	PC7					
e	BR	WO95/34538	12/21/95	PCT	•				
٥	BS	WO96/12021	4/25/96	PCT					
•	ВТ	WO96/14067	5/17/96	PCT					
,	BU	WO97/09346	3/13/97	PCT	•				
,	BV	WO97/09347;	3/13/97	PCT					
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	BBC	WO98/49190	11/5/98	PCT					
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2 30,73	CITATION IN AN APPLICATION		APPLICANT Leland SHAPIRO		
10 ' Jan. E		(PTO-1449)	FILING DATE March 3, 2000	GROUP 1646	
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0	CB ;	Altieri, D.C. J Leukoc Biol 1995, 58	8, <del>120127</del> pp. 120-12	7	
	СС	Anderson ED, Thomas L, Hayflick membrane fusion by a furin-directed 268(33):24887-91 (Nov 25, 1993)			
	CD	Aoki H, Akaike T, Abe K, Kuroda I effect of oryzacystatin, a proteinase in vitro and in vivo. Antimicrob Ag	inhibitor in rice, against her	pes simplex virus type I	
V	CE	Auerswald et al., "K15R M52E) aprotinin is a weak Kunitz-type inhibitor of HIV-replication in H9 cells" <i>Biomed Biochim Acta</i> , 1991, 50(4-6):697-700.			
	CF Ø	Auerswald et al., "Recombinant leech-derived tryptase inhibitor: construction, production, protein chemical characterization and inhibition of HIV-1 replication", Bi Chem Hoppe Seyler, 375(10):695-703 (1994)			
~	CG	Avril, et al., "Identification of the U the V3 loop of HIV-1 gp120 as cath			
J	СН	Avril LE, Di Martino-Ferrer M, Bar associated serine proteinase of U-9 human immunodeficiency virus type 317(1-2):167-72 (Feb 8, 1993)	37 monocytes and peptides fi	rom the V3 loop of the	
	CI	Banfi G, Pirali A, Locatelli M, Mur inhibitor in induced and acquired in infected patients", Scand J Clin Lab	nmunodeficiency. Studies on	transplanted and HIV-	
	CI 🥫	Beck, K.F. et al. in J Exp Biol 1999,	202, 64553 pp. 645-6	653	
	CK P	Bjorck L, Grubb A, Kjellen L. Cysta replication of herpes simplex virus.			
14	CL	Bourinbaiar AS, Lee-Huang S., "Ac guanidinobenzoate, an experimenta Contraception, 51(5):319-22 (May 1)	l vaginal contraceptive with		
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J	CN	Bratt J, Palmblad J. Cytokine-induced endothelial cells. J Immunol 1997 Ju		iury of human	
~	СО	Brinkmann T, Schafers J, Gurtler L, "Inhibition of tryptase TL2 from hun replication of H9 cells by recombina Chem, 16(6):651-60) (Aug 1997)	nan T4+ lymphocytes and in	hibition of HIV-1	
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)	CQ	Bukrinskaia AG, Korneeva MN, Nos reproduction in cultured cells using (Jan-Feb 1989) English Abst	sik DN, Zhdanov VM., "Inh. proteolysis inhibitors", Vop	ibition of HIV r Virusol, 34(1):53-5	
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	<b>9</b> 3	Chesnokova NB, Maichuk YF. Antij Ophthalmol 1986;9(1):593-6	proteases in herpetic keratit	is. Metab Pediatr Syst	
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	CV	Cilberto et al., 1995, Cell, 41:531-54	0		
	CW	Cordiali Fei et al., "Behavior of seve seroconversion period. Comparison 6(2):57-64 (Apr-Jun 1992)			
	сх	Cox et al., "Synergistic combinations immunodeficiency virus", Adv Enzy		on of human	
	CY Ø	Deam DR, Byron KA, Ratnaike S, C antitrypsin phenotypes in homosexuo			
	CZ.	Decroly E, Wouters S, Di Bello C, L "Identification of the paired basic co based on in vitro assays and express 271(48):30442-50 (Nov 1996)	nvertases implicated in HIV	gp 160 processing	
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	CAB	Dery, 0. et al. Am J Physiol 1998, 27	74, C 1429-C 1452		
o .	CAC	DiIanni CL, Drier DA, Deckman IC, I Cordingley MG. Identification of the direct sequence analysis of autoprotect 25;268(3):2048-51	herpes simplex virus-I prot	ease cleavage sites by	
J	CAD	₱ilanni CL, Stevens JT, Bolgar M, O Identification of the serine residue at t protease. J Biol Chem 1994 Apr 29;2	the active site of the herpes		
,	CAE	Ding, A. et al., in J. Immunol. 1990, 1	145, 940		
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	CAG	Flaitz CM, Hicks MJ. "Molecular pir 1998 Nov;34(6):448-53	racy: the viral link to carcine	ogenesis." Oral Oncol	
J	CAH	Franzusoff A, Volpe AM, Josse D, Pi definition of the cellular protease requ 1995 Feb 17;270(7):3154-9	uired for HIV-1 gp160 proc	cessing", J Biol Chem	
~	CAI	Glozman VN, "Immunologic foundati orchiepididymitis", Antibiot Khimiote	ion of enzyme therapy of pa er, 35(7):50-52 ( Jul 1990)a	<sub>tients with</sub> Englisn bstract @ page	
	CAJ	Glynn JM, McElligott DL, Mosier DE cells is blocked by ICE-family proteas Immunol 1996 Oct 1;157(7):2754-275	se inhibition but not by a Fa		
•	CAK	Goureau, 0. et al., in Proc. Natl. Aca	ad. Sci. U.S.A. 1993, 90, 42	276	
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ر	CAM	Gu M, Rappaport J, Leppla SH., "Fur maturation of gp160 of HIV-1", FEBS			
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~	CAU	T4+ lymphocytes immunologically	"A novel membrane-bound serine esterase in human eactive with antibody inhibiting syncytia induced by ation", J Biol Chem., 15:265 (35):21979-85 (Dec			
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7	CA W	Kirkeboen, K.A. and Strand, O.A.	n Acta Anaesthesiol Scand 1	999, 43, 275		
	CAX	Koito A, Hattori T, Murakami T, M neutralizing epitope of human imm sequences with the active site of int (1989)	unodeficiency virus type 1 ha	s homologous amino acid		
	CAY	Langer, R. Nature 1998, 392, 5	pp. 5-10			
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0	CBC	Lowenstein C. J. et al. in Proc. Natl.	Acad. Sci. USA, 1993, 9	<sub>0,9730</sub> - 9734 .	
	CBD	McCall, T.B. et al., in Biochem. Biop			
<b>√</b>	CBE	McNeely TB, Dealy M, Dripps DJ, C leukocyte protease inhibitor: a huma, immunodeficiency virus 1 activity in	n saliva protein exhibiting	anti-human	
J	CBF	Meki AR, Mohey El-Dean ZM. Seru alphal-antitrypsin in scorpion enveno			
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